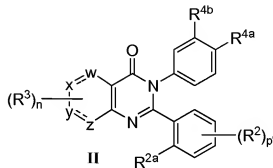


In the claims:

1. Cancelled
2. (Currently amended) The compound according to Claim 1 of the formula II:



wherein a, w, x, y, z, dashed line, R^2 , R^4 , R^6 and R^7 are defined as in Claim 1 for the compound of the Formula I; and

w, x, y and z are independently selected from CH or CH₂;

a dashed line represents an optional double bond;

a is 0 or 1;

b is 0 or 1;

m is 0, 1, or 2;

n is 0 to 2;

r is 0 or 1;

s is 0 or 1;

n is 0 or 1;

p' is 0 to 2;

R^2 is selected from:

- 1) $(C=O)_aC_1-C_{10}$ alkyl,
- 2) $(C=O)_a$ aryl,
- 3) $(C=O)_aNR^6R^7$,
- 4) $(C=O)_aC_3-C_8$ cycloalkyl,
- 5) $(C=O)_a$ heterocyclyl,

- 6) $\text{SO}_2\text{NR}^6\text{R}^7$, and
- 7) $\text{SO}_2\text{C}_1\text{-C}_{10}$ alkyl,

said alkyl, aryl, cycloalkyl, and heterocyclyl is optionally substituted with one or more substituents selected from R^4 ;

R^{2a} is selected from: halogen and $(\text{C}_1\text{-C}_6)\text{alkyl}$; and

R^3 is selected from:

- 1) $(\text{C}=\text{O})_a\text{O}_b\text{C}_1\text{-C}_{10}$ alkyl,
- 2) $(\text{C}=\text{O})_a\text{O}_b\text{aryl}$,
- 3) $(\text{C}=\text{O})_a\text{O}_b\text{C}_2\text{-C}_{10}$ alkenyl,
- 4) $(\text{C}=\text{O})_a\text{O}_b\text{C}_2\text{-C}_{10}$ alkynyl,
- 5) CO_2H ,
- 6) halo,
- 7) OH ,
- 8) $\text{O}_b\text{C}_1\text{-C}_6$ perfluoroalkyl,
- 9) $(\text{C}=\text{O})_a\text{NR}^6\text{R}^7$,
- 10) CN ,
- 11) $(\text{C}=\text{O})_a\text{O}_b\text{C}_3\text{-C}_8$ cycloalkyl,
- 12) $(\text{C}=\text{O})_a\text{O}_b\text{heterocyclyl}$,
- 13) $\text{SO}_2\text{NR}^6\text{R}^7$, and
- 14) $\text{SO}_2\text{C}_1\text{-C}_{10}$ alkyl,

said alkyl, aryl, alkenyl, alkynyl, cycloalkyl, and heterocyclyl is optionally substituted with one or more substituents selected from R^4 ;

R^4 is selected from:

- 1) $(\text{C}=\text{O})_a\text{O}_b\text{C}_1\text{-C}_{10}$ alkyl,
- 2) $(\text{C}=\text{O})_a\text{O}_b\text{aryl}$,
- 3) $\text{C}_2\text{-C}_{10}$ alkenyl,
- 4) $\text{C}_2\text{-C}_{10}$ alkynyl,
- 5) $(\text{C}=\text{O})_a\text{O}_b$ heterocyclyl,
- 6) CO_2H ,
- 7) halo,
- 8) CN ,
- 9) OH ,
- 10) $\text{O}_b\text{C}_1\text{-C}_6$ perfluoroalkyl,

- 11) $O_a(C=O)_bNR^6R^7$,
- 12) oxo,
- 13) CHO,
- 14) $(N=O)R^6R^7$, or
- 15) $(C=O)_aO_bC_3-C_8$ cycloalkyl,
- 16) $SO_2C_1-C_{10}$ alkyl,
- 17) $SO_2NR^6R^7$,

said alkyl, aryl, alkenyl, alkynyl, heterocyclyl, and cycloalkyl optionally substituted with one or more substituents selected from R⁵:

R^{4a} and R^{4b} are independently selected from: hydrogen, halogen and (C₁-C₆)alkyl, provided that at least one is not hydrogen, or

R^{4a} and R^{4b} are combined to form a diradical selected from -CH₂CH₂CH₂CH₂-,
-CH₂CH₂CH₂-, -CH=CH-O- and -CH=CH-N-

R⁵ is selected from:

- 1) $(C=O)_rO_s(C_1-C_{10})$ alkyl,
- 2) $O_r(C_1-C_3)$ perfluoroalkyl,
- 3) (C_0-C_6) alkylene-S(O)_mR^a,
- 4) oxo,
- 5) OH,
- 6) halo,
- 7) CN,
- 8) $(C=O)_rO_s(C_2-C_{10})$ alkenyl,
- 9) $(C=O)_rO_s(C_2-C_{10})$ alkynyl,
- 10) $(C=O)_rO_s(C_3-C_6)$ cycloalkyl,
- 11) $(C=O)_rO_s(C_0-C_6)$ alkylene-aryl,
- 12) $(C=O)_rO_s(C_0-C_6)$ alkylene-heterocyclyl,
- 13) $(C=O)_rO_s(C_0-C_6)$ alkylene-N(R^b)₂,
- 14) C(O)R^a,
- 15) (C_0-C_6) alkylene-CO₂R^a,
- 16) C(O)H,
- 17) (C_0-C_6) alkylene-CO₂H, and
- 18) C(O)N(R^b)₂,

said alkyl, alkenyl, alkynyl, cycloalkyl, aryl, and heterocyclyl is optionally substituted with up to three substituents selected from R^b , OH, (C₁-C₆)alkoxy, halogen, CO₂H, CN, O(C=O)C₁-C₆ alkyl, oxo, and N(R^b)₂;

R⁶ and R⁷ are independently selected from:

- 1) H,
- 2) (C=O)O_pC₁-C₁₀ alkyl,
- 3) (C=O)O_pC₃-C₈ cycloalkyl,
- 4) (C=O)O_paryl,
- 5) (C=O)O_pheterocyclyl,
- 6) C₁-C₁₀ alkyl,
- 7) aryl,
- 8) C₂-C₁₀ alkenyl,
- 9) C₂-C₁₀ alkynyl,
- 10) heterocyclyl,
- 11) C₃-C₈ cycloalkyl,
- 12) SO₂R^a, and
- 13) (C=O)NR^b₂.

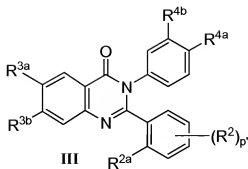
said alkyl, cycloalkyl, aryl, heterocyclyl, alkenyl, and alkynyl is optionally substituted with one or more substituents selected from R⁵, or

R⁶ and R⁷ can be taken together with the nitrogen to which they are attached to form a monocyclic or bicyclic heterocycle with 4-7 members in each ring and optionally containing, in addition to the nitrogen, one or two additional heteroatoms selected from N, O and S, said monocyclic or bicyclic heterocycle optionally substituted with one or more substituents selected from R⁵

R^a is (C₁-C₆)alkyl, (C₃-C₆)cycloalkyl, aryl, or heterocyclyl; and

R^b is H, (C₁-C₆)alkyl, (C₁-C₆)alkyl-NR^a₂, (C₁-C₆)alkyl-NH₂, (C₁-C₆)alkyl-NHR^a, aryl, heterocyclyl, (C₃-C₆)cycloalkyl, (C=O)OC₁-C₆ alkyl, (C=O)C₁-C₆ alkyl or S(O)₂R^a.

3. (Original) A compound of the formula III, or a pharmaceutically acceptable salt or stereoisomer thereof,



wherein

b is 0 or 1;

m is 0, 1 or 2;

p' is 0 to 2;

r is 0 or 1;

s is 0 or 1;

R² is (C₁-C₆)alkylene-NR⁶R⁷; said alkylene is optionally substituted with up to three substituents selected from OH, (C₁-C₆)alkoxy, halogen, CO₂H, CN, O(C=O)C₁-C₆ alkyl, oxo, and NR⁶R⁷;

R^{2a} is selected from: halogen and (C₁-C₆)alkyl;

R^{3a} and R^{3b} are independently selected from: hydrogen and halogen; and

R^{4a} and R^{4b} are independently selected from: hydrogen, halogen, and (C₁-C₆)alkyl, provided that at least one is not hydrogen;

R⁵ is selected from:

- 1) (C=O)_rO_s(C₁-C₁₀)alkyl,
- 2) O_r(C₁-C₃)perfluoroalkyl,
- 3) (C₀-C₆)alkylene-S(O)_mR^a,
- 4) oxo,
- 5) OH,
- 6) halo,
- 7) CN,
- 8) (C=O)_rO_s(C₂-C₁₀)alkenyl,

- 9) $(\text{C}=\text{O})_r\text{O}_8(\text{C}_2\text{-C}_{10})\text{alkynyl}$,
- 10) $(\text{C}=\text{O})_r\text{O}_8(\text{C}_3\text{-C}_6)\text{cycloalkyl}$,
- 11) $(\text{C}=\text{O})_r\text{O}_8(\text{C}_0\text{-C}_6)\text{alkylene-aryl}$,
- 12) $(\text{C}=\text{O})_r\text{O}_8(\text{C}_0\text{-C}_6)\text{alkylene-heterocyclyl}$,
- 13) $(\text{C}=\text{O})_r\text{O}_8(\text{C}_0\text{-C}_6)\text{alkylene-N}(\text{R}^b)_2$,
- 14) $\text{C}(\text{O})\text{R}^a$,
- 15) $(\text{C}_0\text{-C}_6)\text{alkylene-CO}_2\text{R}^a$,
- 16) $\text{C}(\text{O})\text{H}$,
- 17) $(\text{C}_0\text{-C}_6)\text{alkylene-CO}_2\text{H}$, and
- 18) $\text{C}(\text{O})\text{N}(\text{R}^b)_2$,

said alkyl, alkenyl, alkynyl, cycloalkyl, aryl, and heterocyclyl is optionally substituted with up to three substituents selected from R^b , OH, $(\text{C}_1\text{-C}_6)\text{alkoxy}$, halogen, CO_2H , CN, $\text{O}(\text{C}=\text{O})\text{C}_1\text{-C}_6\text{ alkyl}$, oxo, and $\text{N}(\text{R}^b)_2$;

R^6 and R^7 are independently selected from:

- 1) H,
- 2) $(\text{C}=\text{O})\text{O}_b\text{C}_1\text{-C}_{10}\text{ alkyl}$,
- 3) $(\text{C}=\text{O})\text{O}_b\text{C}_3\text{-C}_8\text{ cycloalkyl}$,
- 4) $(\text{C}=\text{O})\text{O}_b\text{aryl}$,
- 5) $(\text{C}=\text{O})\text{O}_b\text{heterocyclyl}$,
- 6) $\text{C}_1\text{-C}_{10}\text{ alkyl}$,
- 7) aryl,
- 8) $\text{C}_2\text{-C}_{10}\text{ alkenyl}$,
- 9) $\text{C}_2\text{-C}_{10}\text{ alkynyl}$,
- 10) heterocyclyl,
- 11) $\text{C}_3\text{-C}_8\text{ cycloalkyl}$,
- 12) SO_2R^a , and
- 13) $(\text{C}=\text{O})\text{NR}^b_2$,

said alkyl, cycloalkyl, aryl, heterocyclyl, alkenyl, and alkynyl is optionally substituted with one or more substituents selected from R^5 , or

R^6 and R^7 can be taken together with the nitrogen to which they are attached to form a monocyclic or bicyclic heterocycle with 4-7 members in each ring and optionally containing, in addition to the nitrogen, one or two additional heteroatoms selected from N, O and S, said monocyclic or bicyclic heterocycle optionally substituted with one or more substituents selected from R^5 ;

R^a is (C₁-C₆)alkyl, (C₃-C₆)cycloalkyl, aryl, or heterocyclyl; and

R^b is H, (C₁-C₆)alkyl, (C₁-C₆)alkyl-NR^a₂, (C₁-C₆)alkyl-NH₂, (C₁-C₆)alkyl-NHR^a, aryl, heterocyclyl, (C₃-C₆)cycloalkyl, (C=O)OC₁-C₆ alkyl, (C=O)C₁-C₆ alkyl or S(O)₂R^a.

4. (Original) The compound according to Claim 3, or the pharmaceutically acceptable salt or stereoisomer thereof, wherein p^{*}, R^{2a}, R^{3a}, R^{3b}, R^{4a}, R^{4b} and R⁵ are as defined for Formula III in Claim 3 and

R² is (C₁-C₆)alkylene-NR⁶R⁷;

R⁶ and R⁷ are independently selected from:

- 1) H,
- 2) C₁-C₁₀ alkyl,
- 3) aryl,
- 4) heterocyclyl,
- 5) C₂-C₁₀ alkenyl,
- 6) C₂-C₁₀ alkynyl, and
- 7) C₃-C₈ cycloalkyl,

said alkyl, cycloalkyl, aryl, heterocyclyl, alkenyl, and alkynyl is optionally substituted with one or more substituents selected from R⁵, or

R⁶ and R⁷ can be taken together with the nitrogen to which they are attached to form a monocyclic or bicyclic heterocycle with 4-7 members in each ring and optionally containing, in addition to the nitrogen, one or two additional heteroatoms selected from N, O and S, said monocyclic or bicyclic heterocycle optionally substituted with one or more substituents selected from R⁵.

5. (Original) A compound selected from:

- 2-(2-bromophenyl)-3-(4-methylphenyl)quinazolin-4(3H)-one;
2-(2-bromophenyl)-3-(4-methylphenyl)-quinazolin-4(3H)-one;
2-(2-chlorophenyl)-3-(4-methylphenyl)-quinazolin-4(3H)-one;
2-(2,4-dichlorophenyl)-3-(4-methylphenyl)quinazolin-4(3H)-one;

2-(2-bromophenyl)-3-(4-chlorophenyl)-quinazolin-4(3H)-one;

2-(2-bromophenyl)-3-(3-fluoro-4-methylphenyl)-quinazolin-4(3H)-one;

3-(3a,7a-dihydro-1H-indol-5-yl)-2-(2-bromophenyl)-quinazolin-4(3H)-one;

6-chloro-2-(2-chlorophenyl)-3-(3-fluoro-4-methylphenyl)-quinazolin-4(3H)-one;

2-(2-chlorophenyl)-3-(3-fluoro-4-methylphenyl)quinazolin-4(3H)-one;

2-(2-methylphenyl)-3-(4-methylphenyl)-quinazolin-4(3H)-one;

7-chloro-2-(2-chlorophenyl)-3-(3-fluoro-4-methylphenyl)quinazolin-4(3H)-one;

2-(2-bromophenyl)-7-chloro-3-(3-fluoro-4-methylphenyl)quinazolin-4(3H)-one;

7-chloro-2-(2-chlorophenyl)-3-(1H-indol-5-yl)quinazolin-4(3H)-one;

2-(2-bromophenyl)-7-chloro-3-(4-chloro-3-fluorophenyl)quinazolin-4(3H)-one;

2-(2-bromophenyl)-3-(3-fluoro-4-methylphenyl)pyrido[2,3-d]pyrimidin-4(3H)-one;

2-(5-bromo-2-chlorophenyl)-7-chloro-3-(3-fluoro-4-methylphenyl)quinazolin-4(3H)-one;

2-(4-bromo-2-chlorophenyl)-7-chloro-3-(3-fluoro-4-methylphenyl)quinazolin-4(3H)-one;

2-(2-chlorophenyl)-3-(3-fluoro-4-methylphenyl)-5,6,7,8-tetrahydroquinazolin-4(3H)-one;

7-chloro-2-{2-chloro-3-[(dimethylamino)methyl]phenyl}-3-(4-chloro-3-fluorophenyl)quinazolin-4(3H)-one ;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-{2-chloro-5-[(4-methylpiperazin-1-yl)methyl]phenyl} quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-{2-chloro-3-[(methylamino)methyl]-phenyl} quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-{2-chloro-3-[(4-methylpiperazin-1-yl)methyl]phenyl} quinazolin-4(3H)-one;

7-chloro-2-{2-chloro-3-[(ethylamino)methyl]phenyl}-3-(4-chloro-3-fluorophenyl)quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-{2-chloro-3-[(isopropylamino)methyl]-phenyl} quinazolin-4(3H)-one;

7-chloro-2-{2-chloro-3-[(cyclobutylamino)methyl]phenyl}-3-(4-chloro-3-fluorophenyl)quinazolin-4(3H)-one;

2-[3-(azetidin-1-ylmethyl)-2-chlorophenyl]-7-chloro-3-(4-chloro-3-fluorophenyl)quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-[2-chloro-3-(pyrrolidin-1-ylmethyl)phenyl]quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-(2-chloro-3-[(3S)-3-hydroxypyrrolidin-1-yl]methyl)phenyl]quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-(2-chloro-3-[(3S)-3-(methoxymethyl)pyrrolidin-1-yl]methyl)phenyl]quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-{2-chloro-3-[(pyrrolidin-3-ylamino)methyl]phenyl}quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-[2-chloro-3-(morpholin-4-ylmethyl)phenyl]quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-[2-chloro-3-(piperidin-1-ylmethyl)phenyl]quinazolin-4(3H)-one;

2-{3-[(4-aminopiperidin-1-yl)methyl]-2-chlorophenyl}-7-chloro-3-(4-chloro-3-fluorophenyl)quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-{2-chloro-3-[(piperidin-4-ylamino)methyl]phenyl}quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-{2-chloro-3-[(4-fluoropiperidin-1-yl)methyl]phenyl}quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-[2-chloro-3-(piperazin-1-ylmethyl)phenyl]quinazolin-4(3H)-one;

2-{3-[(4-acetyl)piperazin-1-yl)methyl]-2-chlorophenyl}-7-chloro-3-(4-chloro-3-fluorophenyl)quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-(2-chloro-3-{{4-(methylsulfonyl)piperazin-1-yl)methyl}phenyl}quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-(2-chloro-3-{{(2-hydroxyethyl)amino}methyl}phenyl}quinazolin-4(3H)-one;

7-chloro-2-[2-chloro-3-({2-(dimethylamino)ethyl}amino)methyl}phenyl]-3-(4-chloro-3-fluorophenyl)quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-(2-chloro-3-{{(2-morpholin-4-ylethyl)amino}methyl}phenyl}quinazolin-4(3H)-one;

2-{3-[(3-aminopyrrolidin-1-yl)methyl]-2-chlorophenyl}-7-chloro-3-(4-chloro-3-fluorophenyl)quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-[2-chloro-3-({(1-methylpiperidin-3-yl)methyl}amino)methyl}phenyl]quinazolin-4(3H)-one;

2-(3-{{3-(aminomethyl)-1-methyl-1 λ 5-piperidin-1-yl}methyl}-2-chlorophenyl)-7-chloro-3-(4-chloro-3-fluorophenyl)quinazolin-4(3H)-one;

2-{3-[(benzylamino)methyl]-2-chlorophenyl}-7-chloro-3-(4-chloro-3-fluorophenyl)quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-{2-chloro-5-[(4-methylpiperazin-1-yl)methyl]phenyl}quinazolin-4(3H)-one;

7-chloro-2-{2-chloro-5-[(ethylamino)methyl]phenyl}-3-(4-chloro-3-fluorophenyl)quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-{2-chloro-5-[(isopropylamino)methyl]phenyl}quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-[2-chloro-5-(pyrrolidin-1-ylmethyl)phenyl]quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-{2-chloro-5-[(pyrrolidin-3-ylamino)methyl]phenyl}quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-[2-chloro-5-(morpholin-4-ylmethyl)phenyl]quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-[2-chloro-5-(piperidin-1-ylmethyl)phenyl]quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-{2-chloro-5-[(piperidin-4-ylamino)methyl]phenyl}quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-[2-chloro-5-(piperazin-1-ylmethyl)phenyl]quinazolin-4(3H)-one;

7-chloro-3-(4-chloro-3-fluorophenyl)-2-(2-chloro-5-{{4-(methylsulfonyl)piperazin-1-yl}methyl}phenyl)quinazolin-4(3H)-one; and

7-chloro-2-[2-chloro-5-({2-(dimethylamino)ethyl}amino)methyl]phenyl]-3-(4-chloro-3-fluorophenyl)quinazolin-4(3H)-one;

or a pharmaceutically acceptable salt thereof.

6. (Currently amended) A pharmaceutical composition that is comprised of a compound in accordance with Claim 42 and a pharmaceutically acceptable carrier.

7. (Original) A pharmaceutical composition that is comprised of a compound in accordance with Claim 3 and a pharmaceutically acceptable carrier.

8. (Currently amended) A method of treating or preventing cancer in a mammal in need of such treatment that is comprised of administering to said mammal a therapeutically effective amount of a compound of Claim 42.

9. Previously cancelled

10. (Original) A method of treating cancer or preventing cancer in accordance with Claim 8 wherein the cancer is selected from cancers of the brain, genitourinary tract, lymphatic system, stomach, larynx and lung.

11. (Original) A method of treating or preventing cancer in accordance with Claim 8 wherein the cancer is selected from histiocytic lymphoma, lung adenocarcinoma, small cell lung cancers, pancreatic cancer, glioblastomas and breast carcinoma.

12.-20. Previously cancelled

21.-24. Cancelled

25.-28. Previously cancelled

29. Cancelled

30. Previously cancelled

31.-34. Cancelled